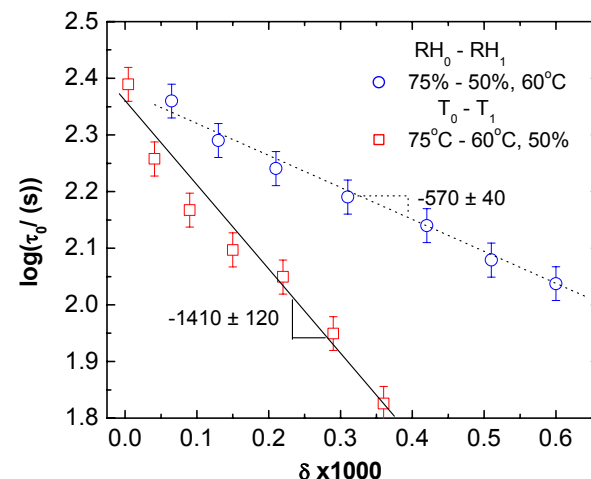


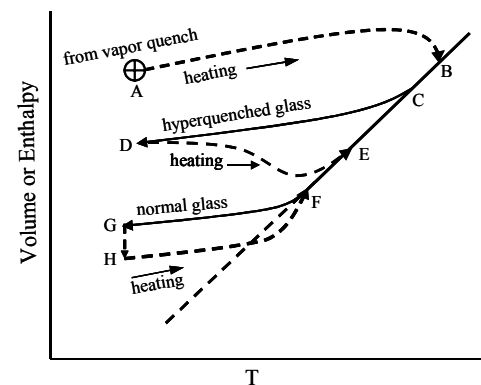
# *The Concentration Glass: A New Glassy State? DMR-0307084*

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- Structural recovery (physical aging) after chemical activity-jumps ( $RH$  or  $PCO_2$ ) is qualitatively similar but quantitatively very different from that after temperature-jumps.*
  - Retardation times are much longer for the same volume departure from equilibrium  $\delta$  (Figure 1)*
- Showed that the “concentration” glass is a different glass from the “temperature” glass. The concentration glass is not equivalent to a temperature-hyper quench formed glass. (Figure 2)*
- First direct demonstration of quantitative differences between concentration- and temperature-glasses.*
- Novel mass uptake measurements during structural recovery are now underway.*
- Preliminary tests of Non-resonant Spectral Hole Burning Dielectric Spectrometer have been performed. Work aims to look at dynamic heterogeneity of concentration glasses*



**Fig. 1.** Plot of creep retardation time vs.  $\delta$  for temperature-glass (red) and concentration-glass (blue) showing that the concentration glass is more stable (retardation times are longer).



**Fig. 2.** Schematic of difference between a temperature hyper-quenched glass and a concentration glass in enthalpy or volume vs. temperature space.

The work in FY 2004 further examined the differences between temperature and concentration glasses. Figure 1 here shows the differences in creep responses as the retardation time for two glasses. The blue data points are for a glass formed from a relative humidity (RH)-jump through the glass transition and the red points show the normal path or temperature-jump through the glass transition. Clearly, for a given volume departure from equilibrium, the concentration glass is “more stable” or has a longer retardation time than does the temperature-glass. In Figure 2 we show an interpretation of our results of volume measurements on a carbon dioxide-created glass when compared with a temperature hyper quenched glass. The volume (or enthalpy) of the PCO<sub>2</sub> glass does not recover towards equilibrium until above the nominal T<sub>g</sub> while the hyper-quenched glass begins changing near to or below the nominal glass transition. The hyper-quench schematic comes from Berens and Hodge, *Macromolecules*, 15, 756 (1982) and we have modified it to make our point about the PCO<sub>2</sub> created glass being different. This figure is in press at *Polymer*.

# *Broad Accomplishments-FY2005*

## *NSF Grant DMR-0307084*

### *• Education and Outreach*

- Project results presented at national and international meetings*
  - Society of Rheology in Oct. 2003.(oral presentations by Y.Zheng, grad. student; M. Alcoutlabi, post-doc)*
  - Rubber Hall of Fame Induction of J.D. Ferry, Nov. 2003 (invited presentation by G.B. McKenna, PI)*
  - American Physical Society in March 2004 (oral presentation by G.B. McKenna, PI)*
  - Society of Plastics Engineers ANTEC in May 2004 (International Award presentation by G.B. McKenna, PI; oral presentation by L. Banda, grad. student)*
  - North American Thermal Analysis Society, October 2004 (L. Banda, grad. Student to give oral presentation.His submitted manuscript was awarded Best Student Paper Award).*
- Project results presented at one regional meeting*
  - NaTex in April 2004 (poster presentations by M. Alcoutlabi, L. Banda, graduate student)*
- Project results presented at U.S. and International Universities and National Labs by PI*
  - NIKE corporation, March 2004 (G.B. McKenna, PI gave invited lecture)*
  - Arizona State University, April 2004 (G.B McKenna, PI gave invited lecture to Chemistry Department)*
  - Colorado School of Mines, April 2004 (G.B. McKenna, PI gave invited lecture to Chem. Engn. Dept.)*
  - Mississippi State University, May 2004 (G.B. McKenna, PI gave invited lecture to Polymer Science Dept.)*
  - University of Messina,Italy, July 2004. (G.B. McKenna, PI gave two invited lectures to graduate students in Physics Department)*

### *• Publication*

- 2 manuscripts published or accepted (J. Polym. Sci., B. Polym.Phys., 42, 2107-2121 (2004); Polymer, in press)*
- 3 proceedings publications*
- 1 manuscript in review (Phys. Rev. Lett.)*